

## INFORMATION CITED BY APPLICANTS THAT MAY BE MATERIAL TO THE PROSECUTION OF THE SUBJECT APPLICATION

Applicants:

W.H. Raskind et al.

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Title:

METHODS FOR IDENTIFYING SUBJECTS SUSCEPTIBLE TO

ATAXIC NEUROLOGICAL DISEASE

## **U.S. PATENT DOCUMENTS**

None

## **FOREIGN PATENT DOCUMENTS**

None

## **OTHER INFORMATION** (Including Author, Title, Date, Pertinent Pages, Etc.)

*Examiner Initial	Cite No.	
DBJ	O1	Abeliovich, A., et al., "PKCy Mutant Mice Exhibit Mild Deficits in Spatial and Contextual Learning," <i>Cell</i> 75:1263-1271, Dec. 31, 1993.
	O2	Brkanac, Z., et al., "A New Dominant Spinocerebellar Ataxia Linked to Chromosome 19q13.4-qter," <i>Arch. Neurol.</i> 59:1291-1295, Aug. 2002.
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DBJ	O4	Burright, E.N., et al., "SCA1 Transgenic Mice: A Model for Neurodegeneration Caused by an Expanded CAG Trinucleotide Repeat," Cell 82(6):937-948, Sept. 22, 1995.
DBJ	O5	Chen, C., et al., "Impaired Motor Coordination Correlates With Persistent Multiple Climbing Fiber Innervation in PKC $\gamma$ Mutant Mice," <i>Cell</i> 83:1233-1242, Dec. 29, 1995.

DBJ	_ 06	Chen, DH., et al., "Cerebral Cavernous Malformation: Novel Mutation in a
		Chinese Family and Evidence for Heterogeneity," J. Neurological Sciences 196:91-96, 2002.
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<del></del>	O16	Quest, A.F.G., et al., "A Phorbol Ester Binding Domain of Protein Kinase Cy: Deletion Analysis of the CYS2 Domain Defines a Minimal 43-Amino Acid Peptide," J. Biol. Chem 269(4):2961-2970, Jan. 28, 1994.
DBJ	O17	Raskind, W.H., et al., "Loss of Heterozygosity in Chondrosarcomas for Markers Linked to Hereditary Multiple Exostoses Loci on Chromosomes 8 and 11," Am. J. Hum. Genet. 56:1132-1139, 1995.

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Examiner

Date Considered

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02/04/2007

TJQ:pt

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